



(19)

Europäisches Patentamt

European Patent Office

Office européen des brevets



(11)

EP 1 009 044 A2

(12)

EUROPEAN PATENT APPLICATION

(43) Date of publication:
14.06.2000 Bulletin 2000/24(51) Int. Cl.⁷: H01L 51/20

(21) Application number: 99203965.1

(22) Date of filing: 25.11.1999

(84) Designated Contracting States:
AT BE CH CY DE DK ES FI FR GB GR IE IT LI LU
MC NL PT SE
Designated Extension States:
AL LT LV MK RO SI

(30) Priority: 09.12.1998 US 208172

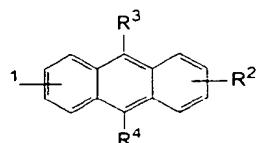
(71) Applicant: EASTMAN KODAK COMPANY
Rochester, New York 14650 (US)

(72) Inventors:
• Shi, Jianmin
Rochester, New York 14650-2201 (US)
• Tang, Ching W.
Rochester, New York 14650-2201 (US)
• Klubek, Kevin P.
Rochester, New York 14650-2201 (US)

(74) Representative:
Lewandowsky, Klaus, Dipl.-Ing. et al
Kodak Aktiengesellschaft,
Patentabteilung
70323 Stuttgart (DE)

(54) Electroluminescent device with anthracene derivatives hole transport layer

(57) An organic multilayer electroluminescent device including an anode and cathode, and including therebetween a hole transport layer; and an electron transport layer disposed in operative relationship with the hole transport layer; wherein the hole transport layer includes an organic compound having the formula:



wherein:

substituents R¹, R², R³ and R⁴ are each individually hydrogen, or alkyl of from 1 to 24 carbon atoms; aryl or substituted aryl of from 5 to 20 carbon atoms; or heteroaryl or substituted heteroaryl of from 5 to 24 carbon atoms; or fluorine, chlorine, bromine; or cyano group.

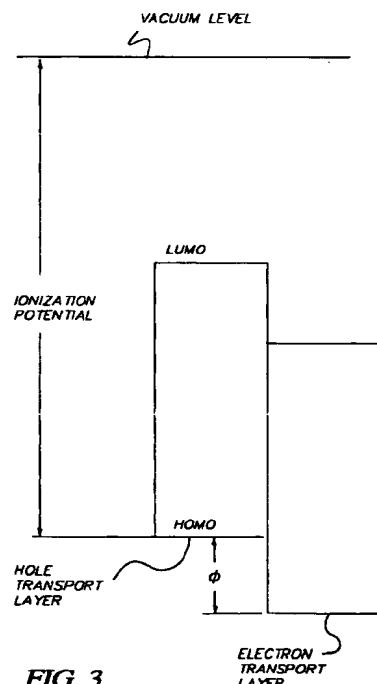


FIG. 3

EP 1 009 044 A2

BEST AVAILABLE COPY

Description

[0001] The present invention relates to organic electroluminescent devices.

[0002]. Organic electroluminescent devices are a class of opto-electronic devices where light emission is produced in response to an electrical current through the device. (For brevity, EL, the common acronym for electroluminescent, is sometimes substituted.) The term organic light emitting diode or OLED is also commonly used to describe an organic EL device where the current-voltage behavior is non-linear, meaning that the current through the EL device is dependent on the polarity of the voltage applied to the EL device. In this embodiment, the term EL and EL devices will include devices described as OLED.

[0003] Organic EL devices generally have a layered structure with an organic luminescent medium sandwiched between an anode and a cathode. The organic luminescent medium usually refers to an organic light emitting material or a mixture thereof in the form of a thin amorphous or crystalline film. Representatives of earlier organic EL devices are disclosed in US-A-3,172,862; US-A-3,173,050; Dresner, "Double Injection Electroluminescence in Anthracene", RCA Review, Vol. 30, pp. 322-334, 1969; and US-A-3,710,167, issued. In these prior arts, the organic luminescent medium was formed of a conjugated organic host material and a conjugated organic activating agent having condensed benzene rings. Naphthalene, anthracene, phenanthrene, pyrene, benzopyrene, chrysene, picene, carbazole, fluorene, biphenyl, terphenyls, quarterphenyls, triphenylene oxide, dihalobiphenyl, trans-stilbene, and 1,4-diphenylbutadiene were offered as examples of organic host materials. Anthracene, tetracene, and pentacene were named as examples of activating agents. The organic luminescent medium was present as a single layer having a thickness much above 1 micrometer. The voltage required to drive the EL devices was as much as a few hundreds volts, thus the luminous efficiency of these EL devices was rather low.

[0004] In commonly-assigned US-A-4,356,429, the art of organic EL device was further advanced by disclosing a bi-layer EL device configuration. The organic luminescent medium in this bi-layer configuration comprises of two extremely thin layers of organic film (<1.0 micrometer in combined thickness) sandwiched between the anode and cathode. The layer adjacent to the anode, termed the hole-transport layer, is specifically chosen to transport predominantly holes only in the EL device. Likewise, the layer adjacent to the cathode is specifically chosen to transport predominantly electrons only in the EL device. The interface or junction between the hole-transport layer and the electron-transport layer is referred to as the electron-hole recombination zone where the electron and hole recombine to produce electroluminescence with the least interference from the electrodes. This recombination zone can be extended beyond the interface region to include portions of the hole-transport layer or the electron-transport layer or both. The extremely thin organic luminescent medium offers reduced electrical resistance, permitting higher current densities for a given voltage applied on the EL device. Since the EL intensity is directly proportional to the current density through the EL device, this thin bi-layer construction of the organic luminescent medium allows the EL device to be operated with a voltage as low as a few volts, in contrast to the earlier EL devices. Thus, the bi-layer organic EL device has achieved a high luminous efficiency in terms of EL output per electrical power input and is therefore useful for applications such as flat-panel displays and lighting.

[0005] Commonly-assigned US-A-4,356,429 disclosed an EL device formed of an organic luminescent medium including a hole transport layer containing 1000 Angstroms of a porphyrinic compound such as copper phthalocyanine, and an electron transport layer of 1000 Angstroms tetraphenylbutadiene in poly(styrene). The anode was formed of a conductive indium-tin-oxide (ITO) glass and the cathode was a layer of silver. The EL device emitted blue light when biased at 20 volts at an average current density in the 30 to 40 mA/cm² range. The brightness of the device was 5 cd/m².

[0006] Further improvements in the bi-layer organic EL devices were taught by commonly-assigned US-A-4,539,507. Dramatic improvements were realized in EL luminous efficiency by substituting the porphyrinic compounds of Tang in the hole-transport layer with an amine compound. With an aromatic tertiary amine such as 1,1-bis(4-di-p-tolylaminophenyl)cyclohexane as the hole-transport layer and an electron transport layer of 4,4'-bis(5,7-di-t-pentyl-2-benzoxazolyl)-stilbene, the EL device was capable of emitting blue-green light with a quantum efficiency of about 1.2 % photon per injected charge when biased at about 20 volts.

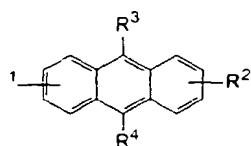
[0007] The use of aromatic amines as the material for the hole-transport layer in organic EL devices has since been generally recognized as numerous prior arts have disclosed the utility of various classes of amines in enhancing the EL device performance. Improvements in the hole-transport material parameters include higher hole transport mobility, more amorphous structures, higher glass transition temperature, and better electrochemical stability. Improvements in the organic EL devices with these improved amines include higher luminous efficiency, longer operational and storage life, and a greater thermal tolerance. For example, the improved arylamine hole transport materials have been disclosed in commonly-assigned US-A-5,061,569. A series of aromatic amines with glass transition temperature as high as 165 °C designed for high temperature EL devices has been disclosed in commonly-assigned US-A-5,554,450. A novel π-conjugated starburst molecule 4,4',4"-tris(3-methylphenylamino) triphenylamine (m-MTDATA), which forms a stable amorphous glass and functions as an excellent hole transport material, was disclosed in US-A-5,374,489.

[0008] The use of organic compounds outside the aromatic amines class for the hole-transport layer in organic EL devices is not common, given the well-known hole-transport properties of the aromatic amines. However, there is a significant disadvantage of using aromatic amines as the hole-transport layer in the bi-layer EL device. Since amines are generally strong electron donors, they can interact with the emissive materials used in the electron-transport layer, resulting in the formation of fluorescence quenching centers and a reduction in the EL luminous efficiency.

[0009] It is an object of the present invention to provide organic compounds outside the class of aromatic amines as the hole transport layer in organic EL devices, which result in enhanced EL performance.

[0010] This object is achieved in an organic multilayer electroluminescent device including an anode and cathode, and comprising therebetween:

10 a hole transport layer; and
an electron transport layer disposed in operative relationship with the hole transport layer;
wherein:
the hole transport layer includes an organic compound having formula I:



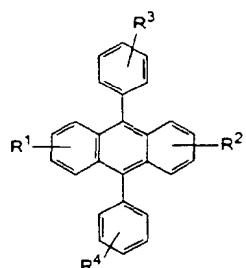
I

wherein:

25 substituents R¹, R², R³ and R⁴ are each individually hydrogen, or alkyl of from 1 to 24 carbon atoms; aryl or substituted aryl of from 5 to 20 carbon atoms; or heteroaryl or substituted heteroaryl of from 5 to 24 carbon atoms; or fluorine, chlorine, bromine; or cyano group.

[0011] Representative examples of the hole transport layer material include:

30 a) Anthracene derivatives having formula II:

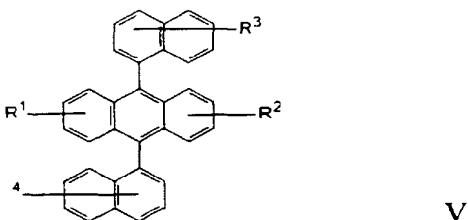
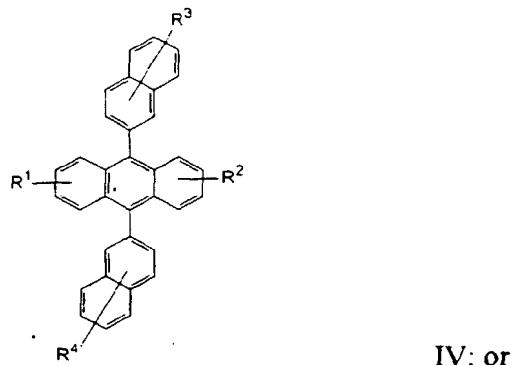
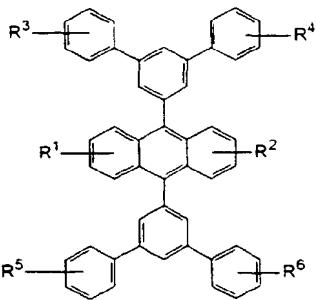


II

45 wherein:

substituents R¹, R², R³ and R⁴ are each individually hydrogen, or alkyl of from 1 to 24 carbon atoms; aryl or substituted aryl of from 5 to 20 carbon atoms; or heteroaryl or substituted heteroaryl of from 5 to 24 carbon atoms; or fluorine, chlorine, bromine; or cyano group;

50 b) Anthracene derivatives having formulas III, IV, V:

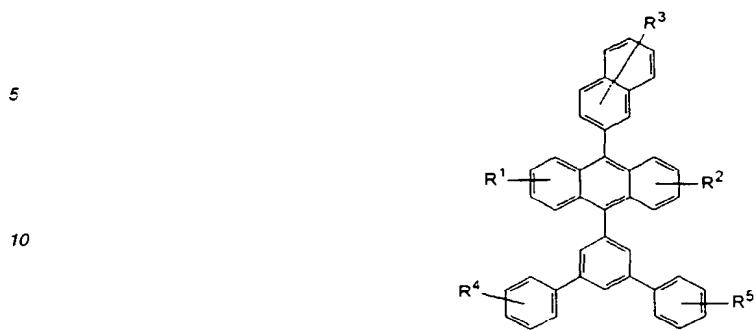


45 wherein:

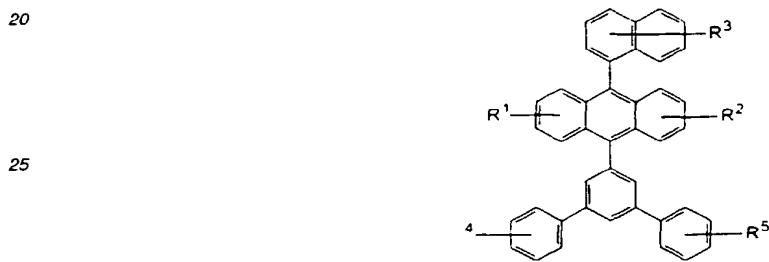
50 substituents R¹, R², R³, R⁴, R⁵ and R⁶ are each individually hydrogen, or alkyl of from 1 to 24 carbon atoms; aryl or substituted aryl of from 5 to 20 carbon atoms; or heteroaryl or substituted heteroaryl of from 5 to 24 carbon atoms; or fluorine, chlorine, bromine; or cyano group; and

c) Anthracene derivatives having formulas VI, VII VIII, IX, X and XI:

55



VI;



VII;

35

40

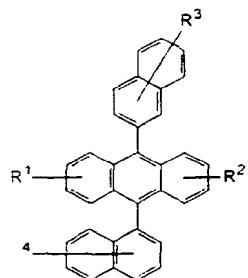
45

50

55

5

5

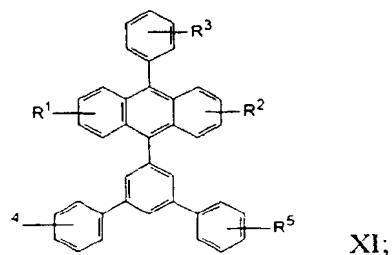


10

15

20

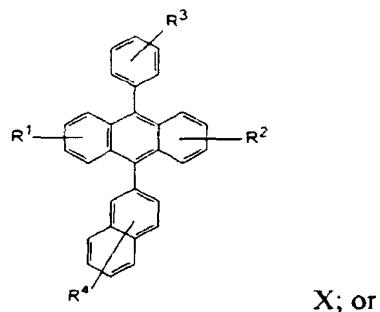
25



30

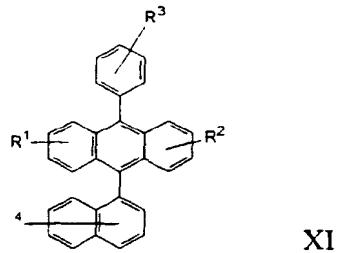
35

40



45

50



55 wherein:

substituents R¹, R², R³, R⁴ and R⁵ are each individually hydrogen, or alkyl of from 1 to 24 carbon atoms; aryl or substituted aryl of from 5 to 20 carbon atoms; or heteroaryl or substituted heteroaryl of from 5 to 24 carbon

atoms; or fluorine, chlorine, bromine; or cyano group.

[0012] Aromatic hydrocarbons or fused aromatic hydrocarbons that are used in the hole transporting layer have the feature that they do not need to include alkylamino- or arylamino- moieties;

5 [0013] The organic compounds in accordance with the present invention have an ionization potential larger than 5.0 eV.

[0014] The hole transport layer in accordance with the present invention effectively works with the electron transport layer or an emissive layer or an electron transport layer which also functions as an emissive layer to provide a highly efficient electroluminescent device.

10 FIG. 1 illustrates the cross-section of a bi-layer organic EL device;
 FIG. 2 illustrates the cross-section of an EL device with a modified bi-layer structure; and
 FIG. 3 illustrates the energy level diagram of an organic EL device with a bi-layer structure as described in FIG. 1.

15 [0015] FIG. 1 illustrates the basic structure used in the construction of organic EL device of this invention. It is a bi-layer structure comprising an organic hole transport layer 30 and an organic electron transport layer 40. The electron transport layer is also the emissive layer from which electroluminescence originates. Together, they form the organic EL medium 50. The anode 20 is adjacent to the hole transport layer and the cathode 60 is adjacent to the electrode transport layer. The substrate is layer 10. This figure is for illustration only and the individual layer thickness is not scaled according to the actual thickness.

20 [0016] FIG. 2 illustrates an alternative construction of the organic EL device of this invention. It is a modified bi-layer structure. The EL medium contains an emissive layer between the hole transport layer and the electron transport layer. This emissive layer is the layer from which electroluminescence originates. Thus, layer 300 is the hole transport layer, layer 400 is the emissive layer, layer 500 is the electron transport layer, and together they form the electroluminescent 25 medium 600. Layer 200 is the anode and layer 700 is the cathode. The substrate is layer 100. This figure is for illustration only and the individual layer thickness is not scaled according to the actual thickness.

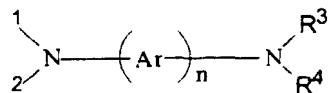
30 [0017] FIG. 3 illustrates the energy level diagram of an organic EL device with a bi-layer structure as described in FIG. 1. The organic EL medium is represented by a hole-transport layer with a characteristic low ionization potential energy, and an electron transport layer with a relatively higher ionization potential energy. The ionization potential energy or ionization potential (IP) for a molecular solid is defined as the energy difference between the vacuum level and the highest occupied molecular orbital (HOMO) level of the solid. The vacuum level is usually referred to as the reference level from which the energy levels of the molecular solid are measured. The HOMO is the highest energy level filled with an electron and in which the hole is free to move. Similarly, the lowest occupied molecular orbital (LUMO) is the lowest energy level devoid of an electron and in which a free electron is free to move. The energy difference 35 between HOMO and LUMO is the bandgap within which there are no available molecular orbital states. The IP value is a measure of the minimum energy required to remove an electron from the molecular solid and can be easily obtained experimentally by photoemission techniques which have been well described in the literature.

40 [0018] The bi-layer structure as illustrated in FIG. 1 is designed to confine the electron-hole recombination at the interface between the hole transport layer and the electron transport layer. This confinement is accomplished by establishing either an electron injection barrier or a hole injection barrier or both at the interface. Referring to the hole injection barrier, it is the difference between the HOMO levels of the hole transport and electron transport layers, as indicated by the symbol, ϕ , in FIG. 1. For large ϕ values, > 0.5 eV, the hole migrating through the hole transport layer towards the interface will be unable to overcome the potential energy barrier and will thus be trapped at the hole transport layer side of the interface. Likewise, the electron injection barrier is the difference between the LUMO levels and a large electron 45 injection barrier for electron injection will localize the electron at the electron transport layer side of the interface. As a result of these charge localizations created by a proper choice of the hole transport and electron transport materials, the electron hole pair will tend to recombine at the interface producing electroluminescence which is characteristics of the interface.

50 [0019] Conventional hole transport materials used in EL devices are mostly arylamines because their hole mobility is among the highest found in common organic materials. Materials with a high mobility are desirable for current-driven devices such as organic EL as the voltage required to operate the device will be low. The arylamines are also known to have the lowest ionization potentials among organic materials. Thus, for creating a hole injection barrier between the hole transport layer and the electron transport layer in a bi-layer EL device, arylamines are appropriate. Highly efficient EL devices have been produced using a variety of arylamines as the hole transport layer.

55 [0020] A class of arylamines found particularly useful in organic EL devices is represented by formula VII:

5



10

VII

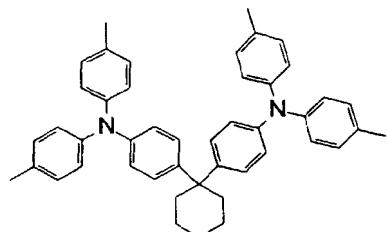
wherein

15

Ar is an arylene group, and arylene moieties are preferably phenyl and phenylene moieties;
 n is an integer of from 1 to 4; and
 R¹, R², R³ and R⁴ are independently selected aryl groups.

20 [0021] These arylamines are particularly useful as the hole transport material in EL devices.

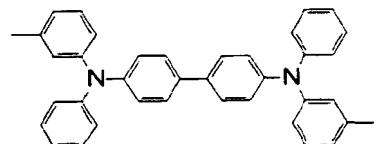
25



30

Compound 1

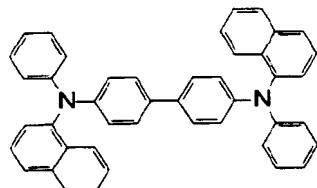
35



40

Compound 2

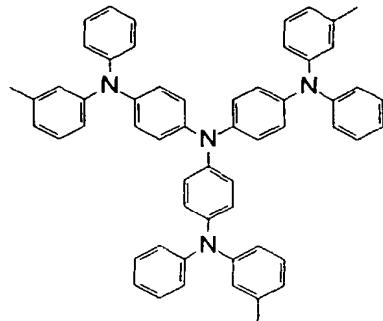
45



50

Compound 3

55



Compound 4

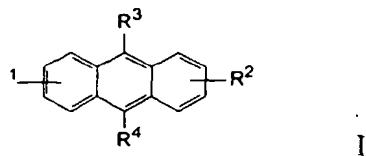
15

[0022] Although arylamines are useful as hole transport materials in EL devices, they do have a number of deficiencies. First, as a class of organic materials, they are relatively strong electron donors, meaning that they can be easily oxidized and therefore are unstable in ambient environments. Second, when used as a hole transport layer adjacent to an electron transport layer in an EL device, the arylamines may interact with the electron transport layer to produce non-emissive centers which will result in a loss of electroluminescence. Third, because of the low ionization potential of the arylamines, the hole injection barrier formed between the arylamine hole transport layer and the electron transport layer will cause the holes to localize in the arylamines which will also result in a loss of electroluminescence. For these reasons, new hole transport materials are useful to further improve the EL device performance.

[0023] The new hole transport materials in this invention include aromatic hydrocarbons or fused aromatic hydrocarbons with a molecular structure containing at least 20 carbon atoms;

[0024] The hole transport layer includes an organic compound having the formula I:

30



wherein:

40 substituents R¹, R², R³ and R⁴ are each individually hydrogen, or alkyl of from 1 to 24 carbon atoms; aryl or substituted aryl of from 5 to 20 carbon atoms; or heteroaryl or substituted heteroaryl of from 5 to 24 carbon atoms; or fluorine, chlorine, bromine; or cyano group.

[0025] Representative examples of the hole transport layer material include:

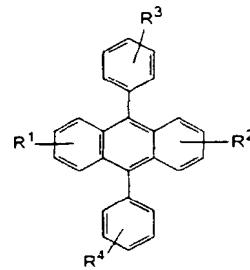
45

a) Anthracene derivatives of formula I:

50

55

5



10

wherein:

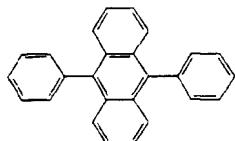
15

substituents R¹, R², R³ and R⁴ are each individually hydrogen, or alkyl of from 1 to 24 carbon atoms; aryl or substituted aryl of from 5 to 20 carbon atoms; or heteroaryl or substituted heteroaryl of from 5 to 24 carbon atoms; or fluorine, chlorine, bromine; or cyano group;

20

The following molecular structures constitute specific examples of anthracene derivatives represented by the general formula II. These compounds are particularly useful as the hole transport material in EL devices.

25



30

Compound 5

35

40

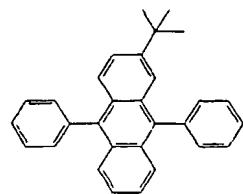
45

50

55

10

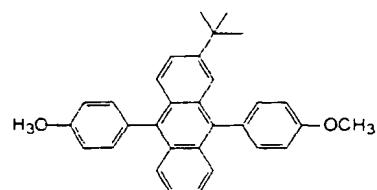
5



Compound 6

10

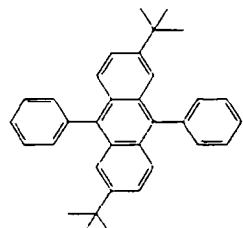
15



Compound 7

20

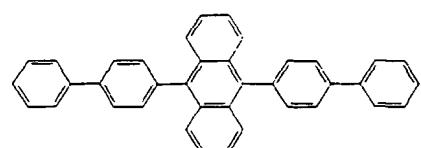
25



Compound 8

30

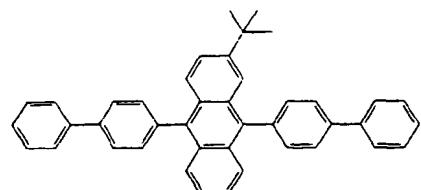
35



Compound 9

40

45

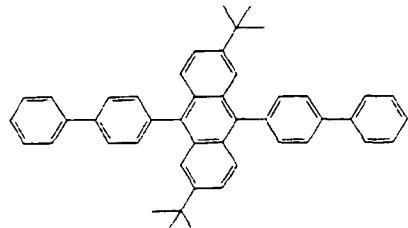


Compound 10

50

55

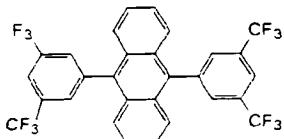
5



10

Compound 11

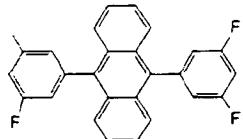
15



20

Compound 12

25



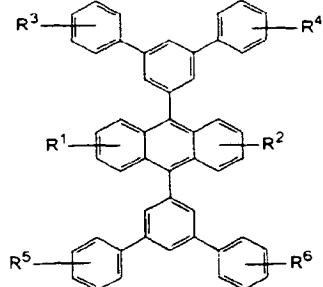
30

Compound 13

b) Anthracene derivatives of formula III, IV, V:

35

40



45

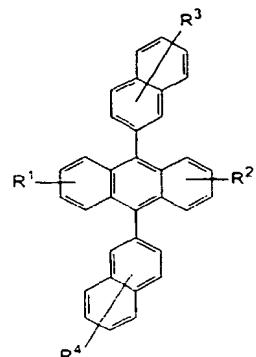
III

50

55

12

5

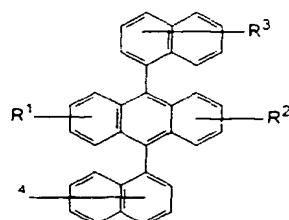


10

15

20

25



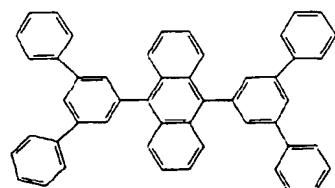
wherein:

30 substituents R¹, R², R³, R⁴, R⁵ and R⁶ are each individually hydrogen, or alkyl of from 1 to 24 carbon atoms; aryl or substituted aryl of from 5 to 20 carbon atoms; or heteroaryl or substituted heteroaryl of from 5 to 24 carbon atoms; or fluorine, chlorine, bromine; or cyano group;

35 The following molecular structures constitute specific examples of anthracene derivatives represented by the general formula III, IV, and V. These compounds are particularly useful as the hole transport material in EL devices.

40

45

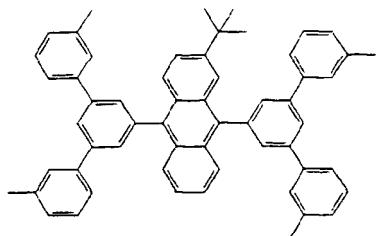


Compound 14

50

55

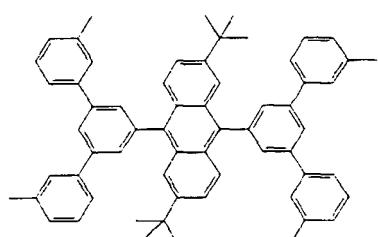
5



10

Compound 15

15

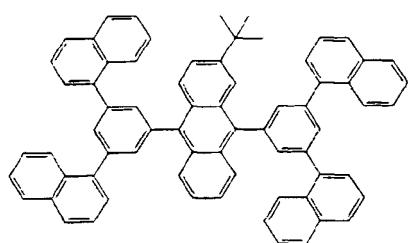


20

Compound 16

25

30

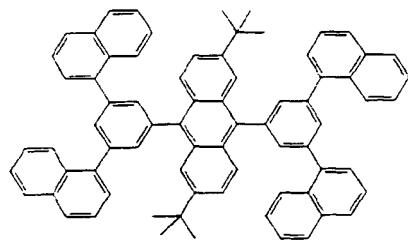


35

Compound 17

40

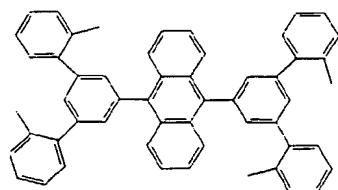
45



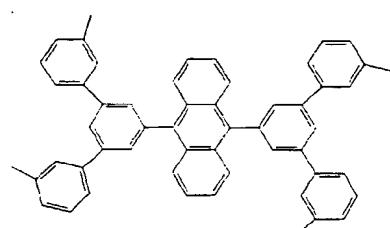
Compound 18

50

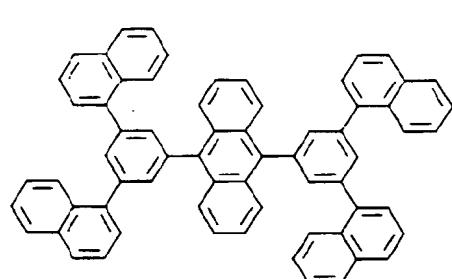
55



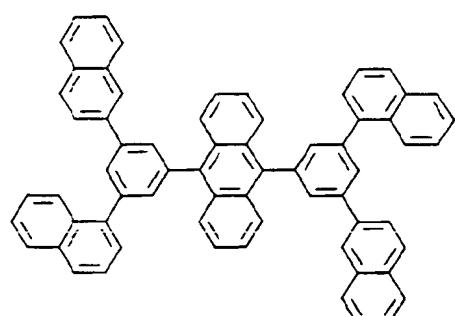
Compound 19



Compound 20



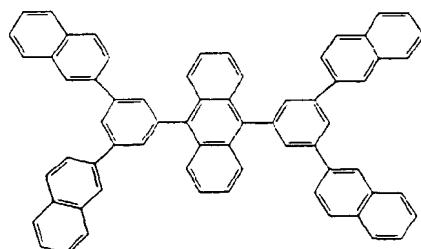
Compound 21



Compound 22

55

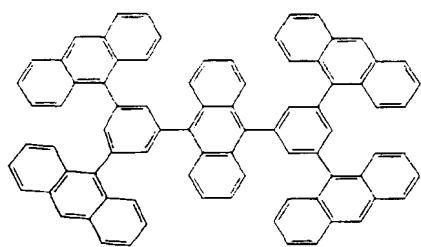
5



10

Compound 23

15

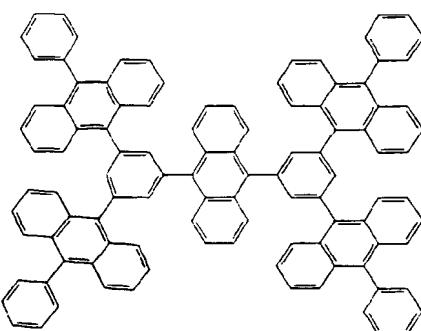


20

Compound 24

25

30

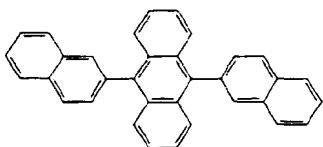


35

Compound 25

40

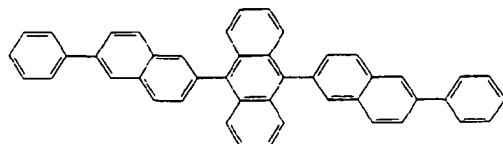
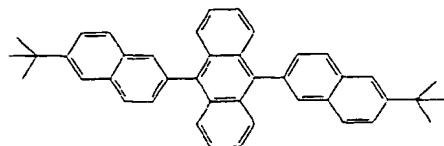
45



Compound 26

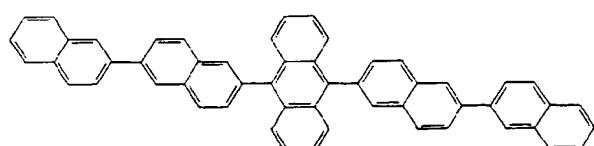
50

55



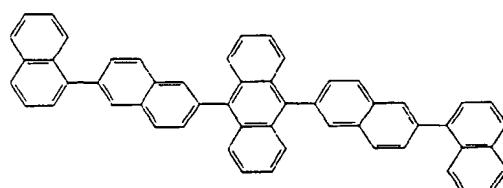
Compound 28

20



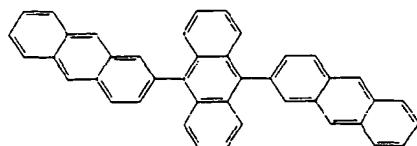
Compound 29

30



Compound 30

40

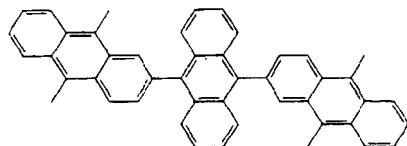


Compound 31

50

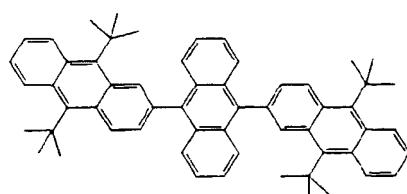
55

5



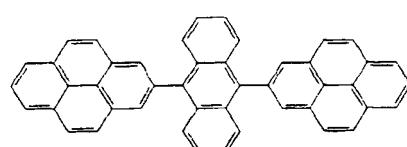
Compound 32

10



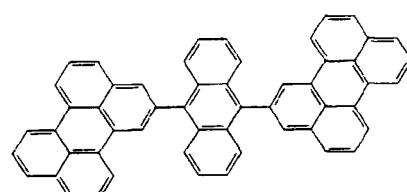
Compound 33

20



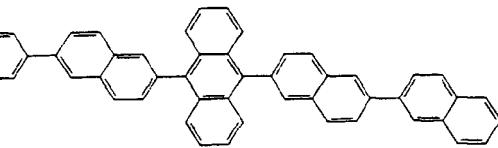
Compound 34

25



Compound 35

30



Compound 36

35

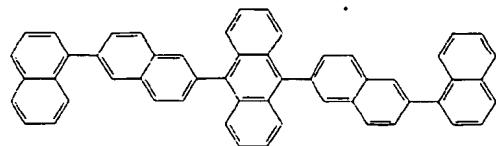
40

45

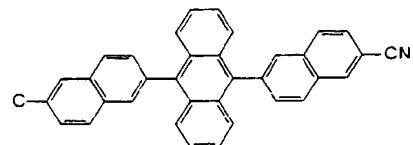
50

55

5



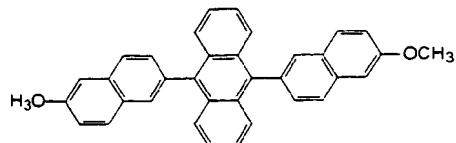
5



Compound 42

10

15

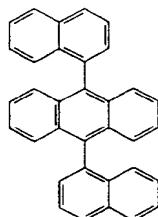


Compound 43

20

25

30

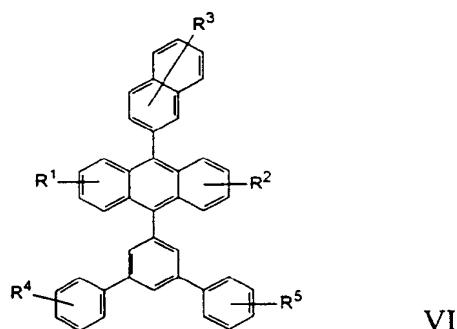


Compound 44

35 c) Anthracene derivatives having formulas VI, VII VIII, IX, X and XI:

40

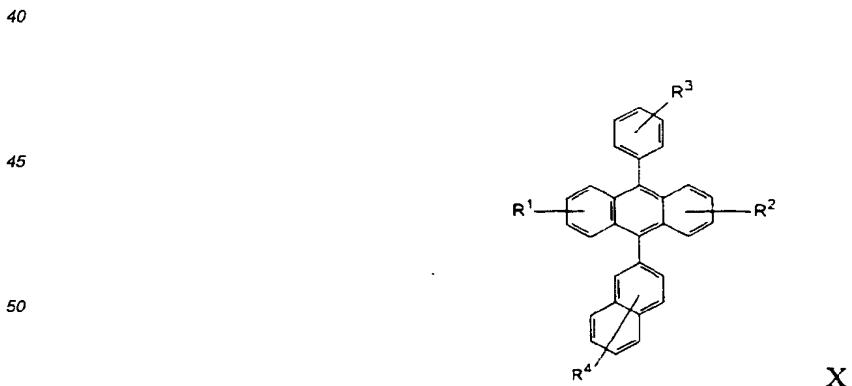
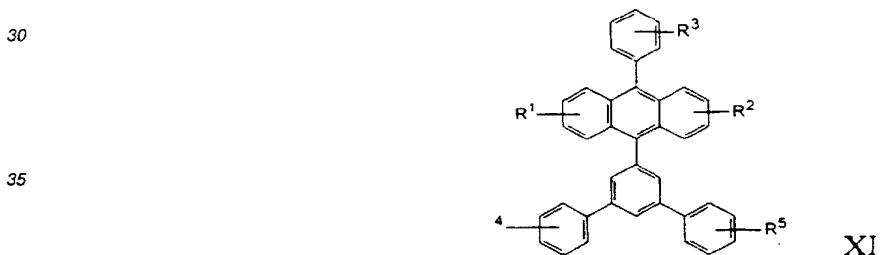
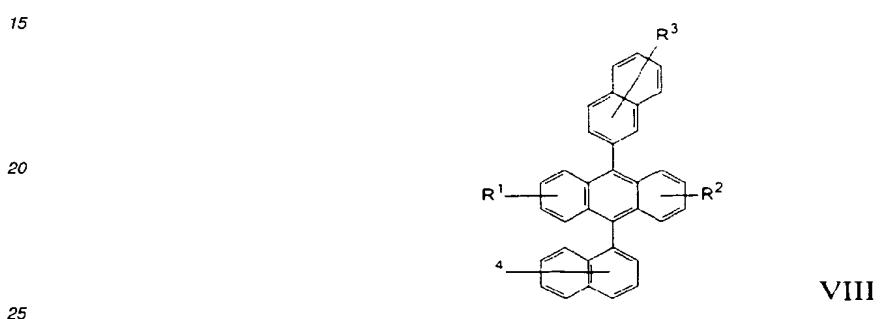
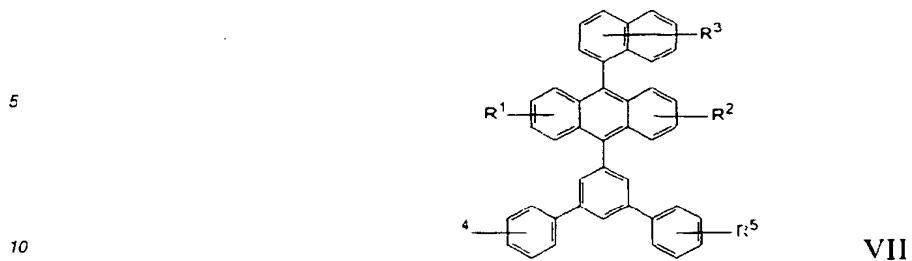
45

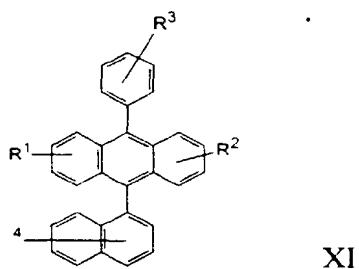


VI

50

55





wherein:

15 substituents R¹, R², R³, R⁴ and R⁵ are each individually hydrogen, or alkyl of from 1 to 24 carbon atoms; aryl or substituted aryl of from 5 to 20 carbon atoms; or heteroaryl or substituted heteroaryl of from 5 to 24 carbon atoms; or fluorine, chlorine, bromine; or cyano group.

20 [0026] The following molecular structures constitute specific examples of anthracene derivatives represented by the general formula VI, VII VIII, IX, X and XI. These compounds are particularly useful as the hole transport material in EL devices.



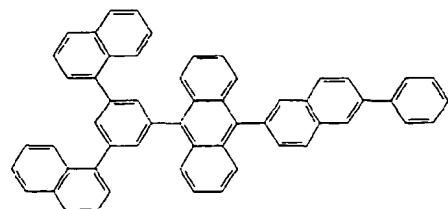
Compound 45



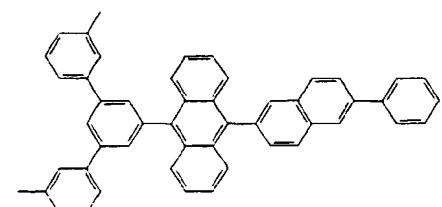
Compound 46

50

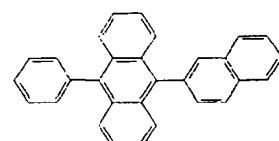
55



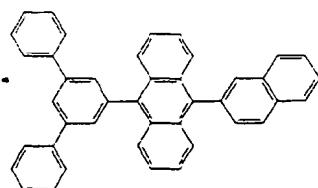
Compound 47



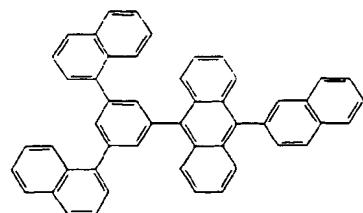
Compound 48



Compound 49



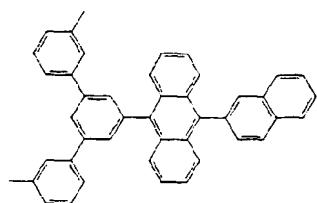
Compound 50



Compound 51

55

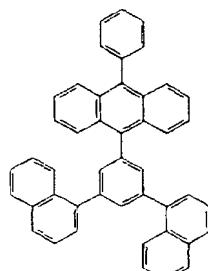
5



10

Compound 52

15

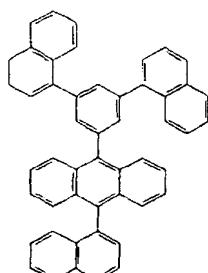


20

Compound 53

25

30

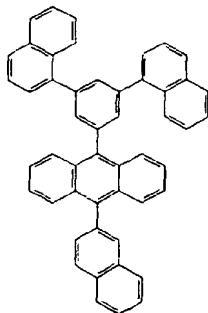


35

Compound 54

40

45

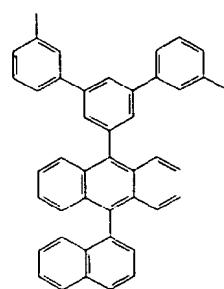


50

Compound 55

55

5



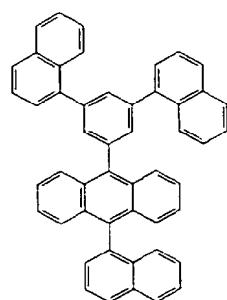
10

Compound 56

15

20

25

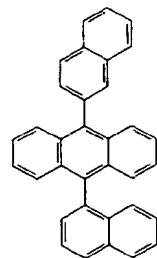


Compound 57

30

35

40



Compound 58

45

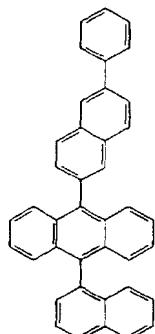
50

55

5

10

15



Compound 59

[0027] The ionization potentials of some of these aromatic hydrocarbon hole transport materials have been measured and their values are compared with the arylamine hole transport materials as follows. It is noted that the aromatic hydrocarbon hole transport materials generally have a higher ionization potential than the arylamines:

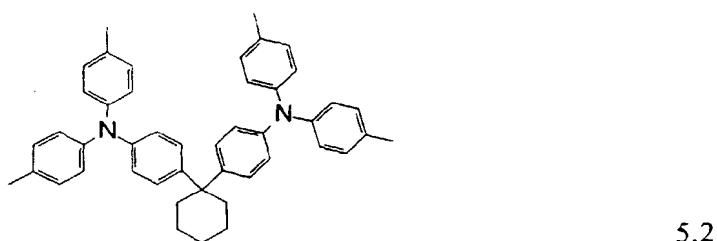
Arylamines and aromatic hydrocarbons IP (eV)

[0028]

25

30

35



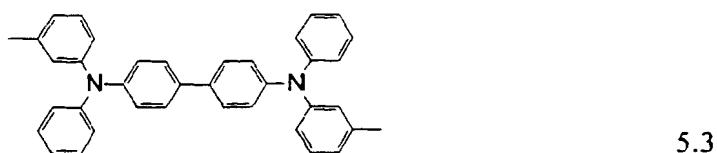
Compound 1

40

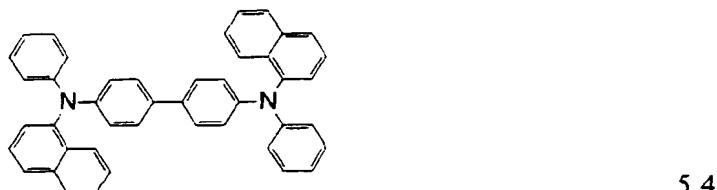
45

50

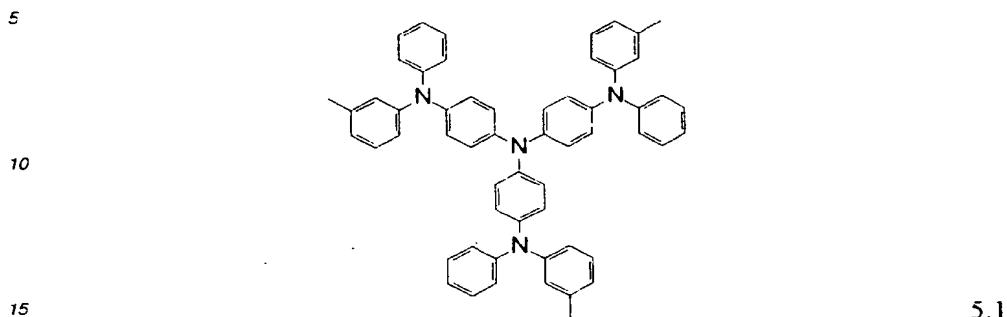
55



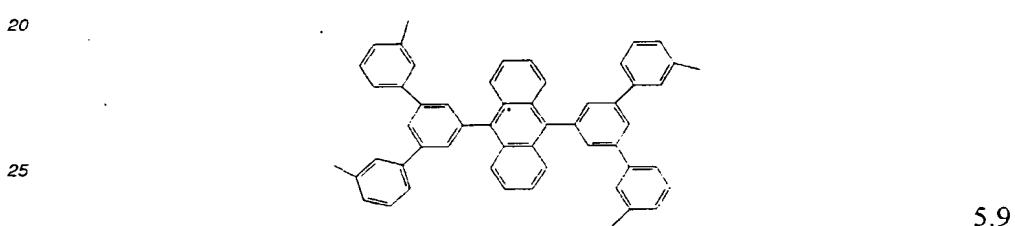
Compound 2



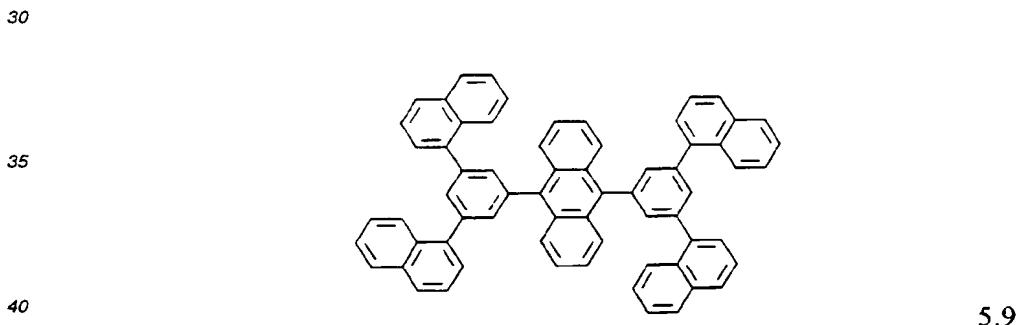
Compound 3



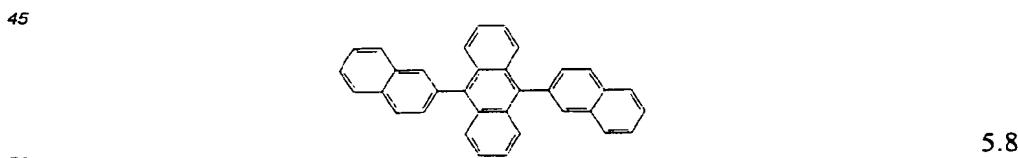
Compound 4



Compound 20



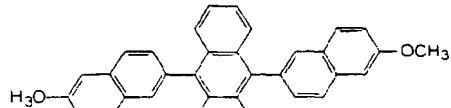
Compound 21



Compound 26

55

5



5.8

10

[0029] A higher ionization potential for the hole transport material is more favorable for hole injection from the hole transport layer to the electron transport layer in a bi-layer EL device resulting in a lower hole injection barrier and consequently a higher EL luminous efficiency. A preferred range for the ionization potentials is 5.0 eV or higher. An alternative criteria is that it should be as high as the ionization potential of the electron transport material in the bi-layer EL device.

[0030] In forming the hole transport layer of the organic EL device, the hole transport materials of this invention can be deposited by a number of methods. The preferred method is by vacuum vapor deposition as these aromatic hydrocarbons have good thermal stability and can be sublimed into thin film. Alternately, they can be dissolved in appropriate solvents and be cast into thin film. Other deposition methods such as printing by the inkjet method, thermal transfer, laser abrasion and sputtering are useful.

[0031] The bi-layer EL device is the basic structure providing high luminous efficiencies and low-voltage operation. Alternative EL device structures have been demonstrated providing improved device performance. These alternative device structures include features in addition to the basic bi-layer structure such as the following structure (a) hole injection layer as disclosed in US-A-4,356,429; (b) cathode modification with alkaline or alkaline halides as disclosed in US-A-5,776,622; (c) anode modification with plasma-deposited fluorochemicals, and (d) doped emitter layer inserted between the hole transport and electron transport layer as disclosed in US-A-4,769,292. These EL device structures retain the hole transport layer as one component of the electroluminescent medium. Therefore, the aromatic hydrocarbon or fused hydrocarbon hole transport materials disclosed in this invention are applicable to these EL device structures as well.

[0032] A preferred EL device structure comprises an anode, a hole transport layer, an emissive layer, and an electron transport layer. In this preferred EL structure, the emissive layer is capable of transporting electrons as well, thus it can be considered as an electron transport layer with the added function of being highly luminescent. The principle function is to provide efficient emissive centers for electroluminescence. This emissive layer comprises a host material doped with one or more fluorescent dyes (FD). The fluorescent dye is usually present in an amount on the order of a few molar percent or less of the host material and it is sufficient to cause the EL emission to be predominantly that of the fluorescent dye. Using this method, highly efficient EL devices can be constructed. Simultaneously, the color of the EL devices can be tuned by using fluorescent dyes of different emission wavelengths. By using a mixture of fluorescent dyes, EL color characteristics of the combined spectra of the individual fluorescent dyes are produced. This dopant scheme has been described in considerable details for EL devices in commonly-assigned US-A-4,769,292.

[0033] An important relationship for choosing a fluorescent dye as a dopant capable of modifying the hue of light emission when present in a host material is a comparison of their bandgap potential which is defined as the energy difference between the highest occupied molecular orbital and the lowest unoccupied molecular orbital of the molecule.

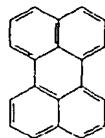
[0034] Preferred host materials for the emissive layer of the organic EL device disclosed in this invention are metal chelated oxinoid compounds, including chelates of oxine itself (also commonly referred to as 8-quinolinol or 8-hydroxy-quinoline or Alq). Another class of preferred host materials is mix ligand 8-quinolinolato aluminum chelates which have been disclosed in US-A-5,141,671. Another class of preferred host materials is distyrylstilbene derivatives as disclosed in US-A-5,366,811.

[0035] For efficient energy transfer from the host to the dopant molecule, a necessary condition is that the band gap of the dopant is smaller than that of the host material. Preferred fluorescent dyes used as the dopant in the emissive layer include coumarins, stilbenes, distyrylstilbenes, derivatives of anthracene, tetracene, perlyenes, rhodamines, and arylamines.

[0036] The molecular structures of the preferred fluorescent dyes for the emissive layer in the EL device are listed as follows:

55

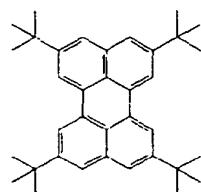
5



10

FD1

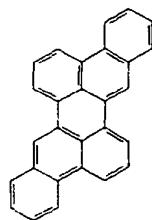
15



20

FD2

25



30

FD3

40

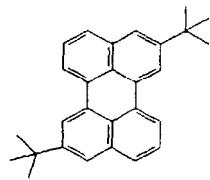
45

50

55

29

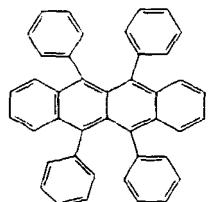
5



10

FD4

15

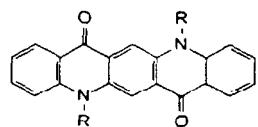


20

FD5

25

30



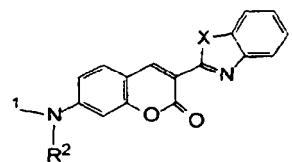
35

FD6 R = H

FD7 R = Me

40

45



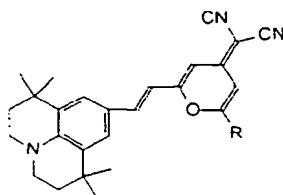
50

FD8 X = O, R¹ = R² = Alkyl

FD9 X = S, R¹ = R² = Alkyl

55

30



FD10	R = Ph
FD11	R = Me
15	FD12 R = t-Bu
20	FD13 R = Mesityl

[0037] Preferred materials for use in forming the electron transporting layer of the organic EL device are metal chelated oxinoid compounds, including chelates of oxine itself (also commonly referred to as 8-quinolinol or 8-hydroxy-quinoline). Representative compounds are 8-hydroquinoline of the group III metals such as Al, In, Mg; and of the group II metals such as Mg, Zn; and of the group I metal such as Li.

[0038] Preferred materials for use in forming the anode of the EL device of this invention are indium tin oxide an anode modified with fluorocarbons.

[0039] Preferred materials for use in forming the cathode of the EL devices of this invention are Mg, Li, or alloys of these materials as disclosed in US-A-5,429,884; and commonly-assigned US-A-5,776,622.

30 Examples

[0040] The invention and its advantages are further illustrated by the specific examples as follows:

35 Example 1 - Synthesis of 3,5-(diphenyl)bromobenzene

[0041] To a solution of 1,3,5-tribromobenzene (60.0 g, 0.19 mol) in 300 mL of dry tetrahydrofuran (THF) was added 0.5 g of bis-(triphenylphosphine)-palladium(II) chloride under nitrogen. After the solution was purged with dry nitrogen for 5 minutes, 175 mL of phenylmagnesium chloride (2.0 M in THF) was added through an addition funnel at room temperature under nitrogen. The reaction mixture was stirred overnight. Then it was quenched by slowly adding 50 mL of 0.5 N HCl with stirring. The solvents were removed via a rotary evaporator. The residue was dissolved in heptane and washed with 0.1 M HCl followed by water. After removal of solvents, crude residues were purified by chromatography on silica gel using 3% methanol in dichloromethane as eluents. After drying, 18.0 g of pure 3,5-(diphenyl)bromobenzene was collected, yielding 30.0%.

45 Example 2 - Synthesis of 9,10-di-(3,5-diphenyl)phenyl anthracene (Compound 14)

[0042] To a suspension of 6.5 g (0.02 mol) of 9, 10-dibromoanthracene and 0.5 g of bis-(triphenylphosphine)-palladium(II) chloride in 100 mL of dry THF at refluxing was added a solution of 3,5-(diphenyl)phenylmagnesium bromide, which was freshly prepared from 15.5 g (0.05 mol) of 3,5-(diphenyl)bromobenzene in 150 mL of dry THF and 1.5 g magnesium in 30 mL of dry THF with 1,2-dibromoethane as an initiator. After addition, the reaction mixture was maintained at reflux for three hours. Then it was cooled and 30 mL of water was carefully added. After removal of the solvents by vacuum rotary evaporator, the residue was extracted by dichloromethane followed by washing with dilute hydrochloric acid and water. The dichloromethane solution was dried over sodium sulfate and passed it through silica gel column. The solvents were removed. The pure 9,10-di-(3',5'-diphenyl)phenyl anthracene (Compound 14) (9.5 g) was obtained by recrystallization from hexane. Yield 75.0 %.

Example 3 - Synthesis of 3,5-di-(m-tolyl)bromobenzene

[0043] To a solution of 1,3,5-tribromobenzene(47.3 g, 0.15 mol) in 150 mL of dry tetrahydrofuran (THF) was added 0.5 g of bis-(triphenylphosphine)-palladium(II) chloride under nitrogen. After the solution was degassed with dry nitrogen for 5 minutes, 155 mL of m-tolyl magnesium bromide (0.2 M in THF) was added through an addition funnel at 70 °C under nitrogen. The reaction mixture was stirred under reflux for another two hours after addition. After cooling the reaction mixture was quenched by slowly adding 50 mL of 0.5 N HCl with stirring. Then the solvents were removed via a rotary evaporator. The residue was dissolved in hexane and washed with 0.1 M HCl followed by water. After removing the solvent, the crude residue was purified by chromatography on silica gel using hexane as the eluent. After drying, 28.0 g of, 3,5-di-m-tolyl bromobenzene was collected. Yield 55.3%.

Example 4 - Synthesis of 9,10-di-(3',5'-m-tolyl)phenyl anthracene (Compound 20)

[0044] To a suspension of 6.5 g (0.02 mol) of 9,10-dibromoanthracene and 0.5 g of bis-(triphenylphosphine)-palladium(II) chloride in 100 mL of dry THF at refluxing was added to a solution of 3,5-di(m-tolyl)phenylmagnesium bromide, which was freshly prepared from 15.5 g (0.046 mol) of 3,5-di-(m-tolyl)bromobenzene in 150 mL of dry THF and clean, dry 1.5 g magnesium in 30 mL of dry THF with 1,2-dibromoethane as an initiator. After the addition, the reaction mixture was kept at reflux for another three hours. Then the reaction mixture was cooled and 30 mL of water was carefully added. After removal of the solvents by a rotary evaporator, the residue was extracted by dichloromethane followed by washing with dilute hydrochloric acid and water. The dichloromethane solution was dried over sodium sulfate and passed through a silica gel column. The pure 9,10-di-(3',5'-m-tolyl)-phenyl anthracene (compound 20) (11.5 g) was obtained by recrystallization from 300 mL of hexane. Yield 76.8%.

Example 5 - Synthesis of 3,5-(1-naphthyl)bromobenzene

[0045] To a solution of 1,3,5-tribromobenzene (105.0 g, 0.22 mol) in 500 mL of dry tetrahydrofuran (THF) was added 1.0 g of bis-(triphenylphosphine)-palladium(II) chloride under nitrogen. After the solution was bubbled with dry nitrogen for 5 minutes, 1-naphthylmagnesium bromide, which was prepared from 150.0 g (0.48 mol) of 1-bromonaphthalene in 100.0 mL of dry THF and clean, dry 18.0 g of magnesium in 250 mL of dry THF with 1,2-dibromoethane as an initiator, was added through an addition funnel at 70 °C under nitrogen. The reaction mixture was stirred under reflux for another two hours. After the reaction mixture was cooled, it was quenched by slowly adding 25.0 mL of 5% HCl with stirring. Then the solvents were removed via a rotary evaporator. The residue was dissolved in dichloromethane and washed with 0.1 M HCl followed by water. After removal of solvents, the crude residue was purified by crystallizing from heptane. A 57.0g of pure 3,5-di(1-naphthyl) bromobenzene was collected. Yield 63.5%.

Example 6 - Synthesis of 9,10-di-[3,5-(1-naphthyl)-phenyl]-anthracene (Compound 21)

[0046] To a suspension of 6.7 g (0.02 mol) of 9,10-dibromoanthracene and 0.3 g of bis-(triphenylphosphine)-palladium(II) chloride in 150 mL of dry THF heated at refluxing was added to a solution of 3,5-di-(1-naphthyl)phenylmagnesium bromide, which was freshly prepared from 18.4 g of (0.045 mol) of 3,5-di-(1-naphthyl)-bromobenzene in 150 mL of dry THF and clean, dry 1.5 g of magnesium in 30 mL of dry THF with 1,2-dibromoethane as an initiator. After the addition, the reaction mixture was kept at reflux for another three hours. Then the reaction mixture was cooled and 30 mL of 0.5% HCl was carefully added. After removal of the solvents by vacuum rotary evaporator, the residue was filtered and washed by water, 1:1 water:acetone, and followed by a minimum amount of dichloromethane. After drying, the pure 9,10-bis-[3',5'-(1-naphthyl)phenyl] anthracene (Compound 21) (12.5 g) was obtained. Yield 74.0%.

Example 7 - Synthesis of 2-naphthylene boronic acid

[0047] A solution of n-BuLi (1.6 M in hexane, 100 mL, 0.16 mol) was added via an addition funnel to 2-bromonaphthalene (30.0 g, 0.14 mol) in 200 mL of dry THF at -78 °C. The yellow suspension was stirred at this temperature for a half hour, a solution of B(OMe)₃ (26.6 mL, 29.1 g, 0.28 mol) in 150 mL of dry THF was added dropwise, with the temperature kept below -60 °C. The resulting colorless solution was allowed to warm to room temperature overnight, then 300 mL of 10 M HCl was added and the mixture stirred for a further one hour under nitrogen. Water and ether were added, and the aqueous layer was extracted several times with ether. The combined organic extracts were dried over MgSO₄ and evaporated under reduced pressure to yield a white solid (21.0 g, 95%), which was used in the coupling reaction without further purification.

Example 8 - Synthesis of 9,10-di-(2-naphthyl)anthracene (Compound 26)

[0048] Pd(PPh₃)₄ (1.0 g, 0.8 mmol) and 300 mL of 2.0 M aqueous Na₂CO₃ were added to a solution of 9,10-dibromoanthracene (34.0 g, 0.1 mol) and 2-naphthylene boronic acid (40.0 g, 0.232 mol) in 600 mL of toluene and 100 mL of ethanol. The reaction mixture was purged with nitrogen for 10 mm. After refluxing under nitrogen overnight, the organic suspension layer was separated while hot and was added 300 mL of 2.0 N HCl and refluxed for one hour with vigorous stirring. The aqueous layer was separated again while hot followed by washing with water three times until pH is about 7. The precipitates from the organic layer was filtered and washed with small amount of cold acetone followed by toluene. After drying, 34.0 g of pure 9,10-di-(2-naphthyl)anthracene (compound 26) was obtained. Yield 80.0%.

Example 9 - Synthesis of 9,10-di-[2-(6-methoxynaphthyl)]anthracene (compound 43)

[0049] To a suspension of 22.0 g (0.09 mol) of 9,10-dibromoanthracene and 0.75 g of bis(triphenylphosphine)palladium(II) chloride in 200 mL of dry THF at reflux was added a solution of 6-methoxy 2-naphthylmagnesium bromide, which was fresh by prepared from 50.0 g (0.211 mol) of 6-methoxy 2-bromonaphthylene in 400 mL of dry THF and 5.6 g of magnesium in 100 mL of dry THF with 1,2-dibromoethane as an initiator. After addition, the reaction mixture was maintained at reflux for three hours. Then it was cooled and 100 mL of THF and 50 mL of 15% hydrochloric acid was carefully added. After removal of the solvents by vacuum rotary evaporator, the residue was filtered and washed with water until pH = 7. The crude product was refluxed in 500 mL of dichloromethane for one hour. After cooling, it was filtered and washed with a small amount of cold acetone to give 34.0 g of pure 9,10-di-[2-(6-methoxynaphthyl)]anthracene (Compound 43). Yield 77.1 %.

EL device fabrication and performanceExamples 10 to 16

[0050] EL devices of this invention were constructed in the following manner. The organic EL medium has an anode, a hole transport layer, an emissive and electron transport layer, and a cathode. The substrate was glass.

- a) The anode was a conductive indium-tin-oxide (ITO) coated on a glass substrate. It was about 1000 Angstroms thick. The ITO glass was cleaned using a commercial glass plate cleaner. Prior to the deposition of the organic layers, the ITO substrate was subjected to an oxygen plasma clean in a commercial etcher.
- b) A hole transport layer about was deposited onto the ITO substrate by vacuum vapor deposition using a tantalum boat source. The layer thickness was about 600 Angstroms.
- c) An electron-transport and emissive layer was deposited on top of the hole transport layer by vacuum vapor deposition using a tantalum boat source. The layer thickness was about 700 Angstroms.
- d) A cathode layer was deposited on top of the electron transport and emissive layer. The layer thickness was about 2000 Angstroms and the atomic composition of the cathode was about 10 parts magnesium and 1 part silver.

[0051] The above deposition sequence was completed in a single sequence without a vacuum break between the deposition of individual layers. The completed EL device was then sealed with a cover glass plate in a dry glove box for protection against ambient environment. A desiccant material was also include in the sealed package to improve the storage life of the EL device.

[0052] The results of the EL devices from examples 10 to 16 are shown in Table I. Example 10 is a comparative example. Compound 3 used in this example is an arylamine. The light output and luminous efficiency obtained by this EL device were substantially lower in comparison with the EL devices of Examples 11 - 16, which used aromatic hydrocarbons as the hole transport layer. An efficiency gain on the order of 30 to 40% has been realized by using an aromatic hydrocarbon as the hole transport layer.

Table I

	Hole transport layer	Electron transport and emissive layer	Applied Voltage (V)	Brightness (cd/m ²)	Efficiency (cd/A)	Color
Example 10	Compound 3	Alq	7.5	578	2.9	Green
Example 11	Compound 26	Alq	6.3	855	4.2	Green

Table I (continued)

	Hole transport layer	Electron trans- port and emis- sive layer	Applied Voltage (V)	Brightness (cd/m ²)	Efficiency (cd/A)	Color	
5	Example 12	Compound 9	Alq	8.6	628	3.14	Green
10	Example 13	Compound 20	Alq	8.9	929	4.65	Green
15	Example 14	Compound 21	Alq	10.7	877	4.22	Green
20	Example 15	Compound 39	Alq	8.6	820	4.10	Green
25	Example 16	Compound 43	Alq	8.5	879	4.39	Green

Examples 17 to 22

[0053] EL devices of this invention were constructed in a manner similar to Examples 10-16. The organic EL medium has an anode, a hole transport layer, an emissive layer, an electron-transport layer, and a cathode. The substrate was glass.

- a) The anode was a conductive indium-tin-oxide (ITO) coated on a glass substrate. It was about 1000 Angstrom thick. The ITO glass was cleaned using a commercial glass plate cleaner. Prior to the deposition of the organic layers, the ITO substrate was subjected to an oxygen plasma clean in a commercial etcher.
- b) A hole transport layer about was deposited onto the ITO substrate by vacuum vapor deposition using a tantalum boat source. The layer thickness was about 600 Angstroms.
- c) An emissive layer was deposited on top of the hole transport layer by vacuum vapor deposition using a tantalum boat source. The layer thickness was about 350 Angstroms.
- d) An electron transport layer was deposited on top of the emissive layer by vacuum vapor deposition using a tantalum boat source. The layer thickness was about 350 Angstroms.
- e) A cathode layer was deposited on top of the electron transport layer. The layer thickness was about 2000 Angstroms and the atomic composition of the cathode was about 10 parts magnesium and 1 part silver.

[0054] The above deposition sequence was completed in a single sequence without a vacuum break between the deposition of individual layers. The completed EL device was then sealed with a cover glass plate in a dry glove box for protection against ambient environment. A desiccant material was also include in the sealed package to improve the storage life of the EL device.

[0055] The results of the EL devices from examples 17 to 22 are shown in Table II. Example 17 is a comparative example using an arylamine (Compound 3) as the hole transport layer. The light output and luminous efficiency obtained from this EL were substantially lower in comparison with the EL device of Example 18, which used an aromatic hydrocarbon of Compound 26 as the hole transport layer instead of an arylamine. Otherwise, both EL devices have an identical structure. An efficiency gain of 34% has been realized in using the aromatic hydrocarbon as the hole transport layer. Example 21 is another comparative example using an arylamine (Compound 3) as the hole transport layer. The red light output and luminous efficiency obtained from this EL were substantially lower in comparison with the EL device of Example 22, which used an aromatic hydrocarbon Compound 26 as the hole transport layer instead of an arylamine. Otherwise, both EL devices have an identical structure. An efficiency gain of 80% has been realized in using the aromatic hydrocarbon as the hole transport layer.

Table II

Examples	Hole trans- port layer	Doped emis- sive layer	Electron transport layer	Applied Volt- age (V)	Brightness (cd/m ²) @ 20 mA/cm ²	Efficiency cd/A	Emitting Light	
50	Example 17	Compound 3	Alq + 1% FD 9	Alq	6.9	2219	11.1	Green
55	Example 18	Compound 26	Alq + 1% FD 9	Alq	6.5	2994	14.9	Green

Table II (continued)

Examples	Hole trans- port layer	Doped emis- sive layer	Electron transport layer	Applied Volt- age (V)	Brightness (cd/m ²) @ 20 mA/cmA ²	Efficiency cd/A	Emitting Light	
5	Example 19	Compound 21	Alq + 1% FD 9	Alq	8.3	3133	15.6	Green
10	Example 20	Compound 43	Alq + 1% FD 9	Alq	8.5	2848	14.24	Green
15	Example 21	Compound 3	Alq+ 1% FD 13	Alq	7.9	439	2.20	Red
	Example 22	Compound 26	Alq+ 1% FD 13	Alq	7.7	791	3.90	Red

PARTS LIST

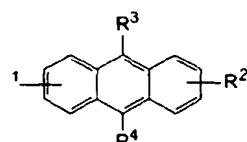
20 [0056]

- 10 substrate
 20 anode
 30 hole transport layer
 25 40 electron transport layer
 50 organic EL medium
 60 cathode
 100 substrate
 200 anode
 300 300 hole transport layer
 400 emissive layer
 500 electron transport layer
 600 EL medium
 700 cathode

35 **Claims**

1. An organic multilayer electroluminescent device including an anode and cathode, and comprising therebetween:
 40 a hole transport layer; and
 an electron transport layer disposed in operative relationship with the hole transport layer;
 wherein:
 the hole transport layer includes an organic compound having the formula:

45



50

wherein:
 substituents R¹, R², R³ and R⁴ are each individually hydrogen, or alkyl of from 1 to 24 carbon atoms; aryl or substituted aryl of from 5 to 20 carbon atoms; or heteroaryl or substituted heteroaryl of from 5 to 24 carbon atoms; or fluorine, chlorine, bromine; or cyano group.

- 55 2. An organic multilayer electroluminescent device including an anode and cathode, and comprising therebetween:

a hole transport layer; and

an electron transport layer disposed in operative relationship with the hole transport layer;

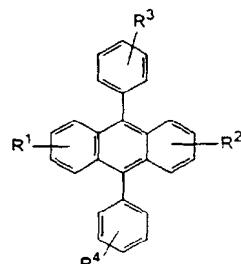
wherein:

the hole transport layer includes an organic compound having the formula:

5

10

15



wherein:

substituents R¹, R², R³ and R⁴ are each individually hydrogen, or alkyl of from 1 to 24 carbon atoms; aryl or substituted aryl of from 5 to 20 carbon atoms; or heteroaryl or substituted heteroaryl of from 5 to 24 carbon atoms; or fluorine, chlorine, bromine; or cyano group.

20 3. An organic multilayer electroluminescent device including an anode and cathode, and comprising therebetween:

25 a hole transport layer; and

an electron transport layer disposed in operative relationship with the hole transport layer;

wherein:

the hole transport layer includes an organic compound having the formula:

30

35

40

45 4. An organic multilayer electroluminescent device including an anode and cathode, and comprising therebetween:

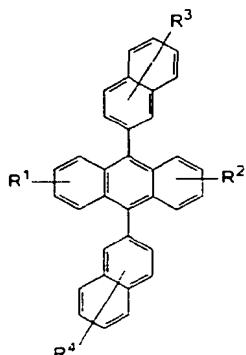
50 a hole transport layer; and

an electron transport layer disposed in operative relationship with the hole transport layer;

wherein:

the hole transport layer includes an organic compound having the formula:

55



wherein:

substituents R¹, R², R³ and R⁴, are each individually hydrogen, or alkyl of from 1 to 24 carbon atoms; aryl or substituted aryl of from 5 to 20 carbon atoms; or heteroaryl or substituted heteroaryl of from 5 to 24 carbon atoms; or fluorine, chlorine, bromine; or cyano group.

20

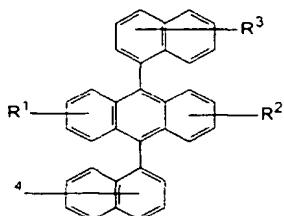
5. An organic multilayer electroluminescent device including an anode and cathode, and comprising therebetween:

a hole transport layer; and

25 an electron transport layer disposed in operative relationship with the hole transport layer;

wherein:

the hole transport layer includes an organic compound having the formula:



wherein:

40 substituents R¹, R², R³ and R⁴ are each individually hydrogen, or alkyl of from 1 to 24 carbon atoms; aryl or substituted aryl of from 5 to 20 carbon atoms; or heteroaryl or substituted heteroaryl of from 5 to 24 carbon atoms; or fluorine, chlorine, bromine; or cyano group.

45

6. An organic multilayer electroluminescent device including an anode and cathode, and comprising therebetween:

a hole transport layer; and

an electron transport layer disposed in operative relationship with the hole transport layer;

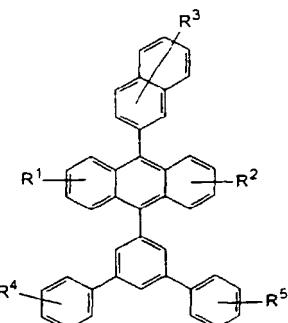
wherein:

the hole transport layer includes an organic compound having the formula:

50

55

5



10

15 wherein:

substituents R¹, R², R³, R⁴ and R⁵ are each individually hydrogen, or alkyl of from 1 to 24 carbon atoms; aryl or substituted aryl of from 5 to 20 carbon atoms; or heteroaryl or substituted heteroaryl of from 5 to 24 carbon atoms; or fluorine, chlorine, bromine; or cyano group.

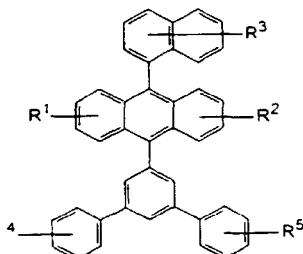
20 7. An organic multilayer electroluminescent device including an anode and cathode, and comprising therebetween:

a hole transport layer; and
an electron transport layer disposed in operative relationship with the hole transport layer;

wherein:

25 the hole transport layer includes an organic compound having the formula:

30



35

40 wherein:

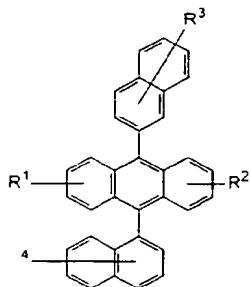
substituents R¹, R², R³, R⁴ and R⁵ are each individually hydrogen, or alkyl of from 1 to 24 carbon atoms; aryl or substituted aryl of from 5 to 20 carbon atoms; or heteroaryl or substituted heteroaryl of from 5 to 24 carbon atoms; or fluorine, chlorine, bromine; or cyano group.

45 8. An organic multilayer electroluminescent device including an anode and cathode, and comprising therebetween:

a hole transport layer; and
an electron transport layer disposed in operative relationship with the hole transport layer;
wherein:
the hole transport layer includes an organic compound having the formula:

50

55



wherein:

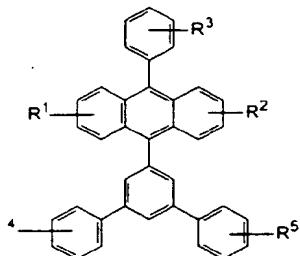
substituents R¹, R², R³, R⁴ and R⁵ are each individually hydrogen, or alkyl of from 1 to 24 carbon atoms; aryl or substituted aryl of from 5 to 20 carbon atoms; or heteroaryl or substituted heteroaryl of from 5 to 24 carbon atoms; or fluorine, chlorine, bromine; or cyano group.

20 9. An organic multilayer electroluminescent device including an anode and cathode, and comprising therebetween:

a hole transport layer; and

an electron transport layer disposed in operative relationship with the hole transport layer; wherein:
the hole transport layer includes an organic compound having the formula:

25



wherein:

substituents R¹, R², R³, R⁴ and R⁵ are each individually hydrogen, or alkyl of from 1 to 24 carbon atoms; aryl or substituted aryl of from 5 to 20 carbon atoms; or heteroaryl or substituted heteroaryl of from 5 to 24 carbon atoms; or fluorine, chlorine, bromine; or cyano group.

40

10. An organic multilayer electroluminescent device including an anode and cathode, and comprising therebetween:

a hole transport layer; and

45

an electron transport layer disposed in operative relationship with the hole transport layer;

wherein:

the hole transport layer includes an organic compound having the formula:

50

55

5

10

15

20 11. An organic multilayer electroluminescent device including an anode and cathode, and comprising therebetween:

a hole transport layer; and
 an electron transport layer disposed in operative relationship with the hole transport layer;
 wherein:

25 the hole transport layer includes an organic compound having the formula:

30

35

wherein:
 substituents R¹, R², R³ and R⁴ are each individually hydrogen, or alkyl of from 1 to 24 carbon atoms; aryl or substituted aryl of from 5 to 20 carbon atoms; or heteroaryl or substituted heteroaryl of from 5 to 24 carbon atoms; or fluorine, chlorine, bromine; or cyano group.

45

50

55

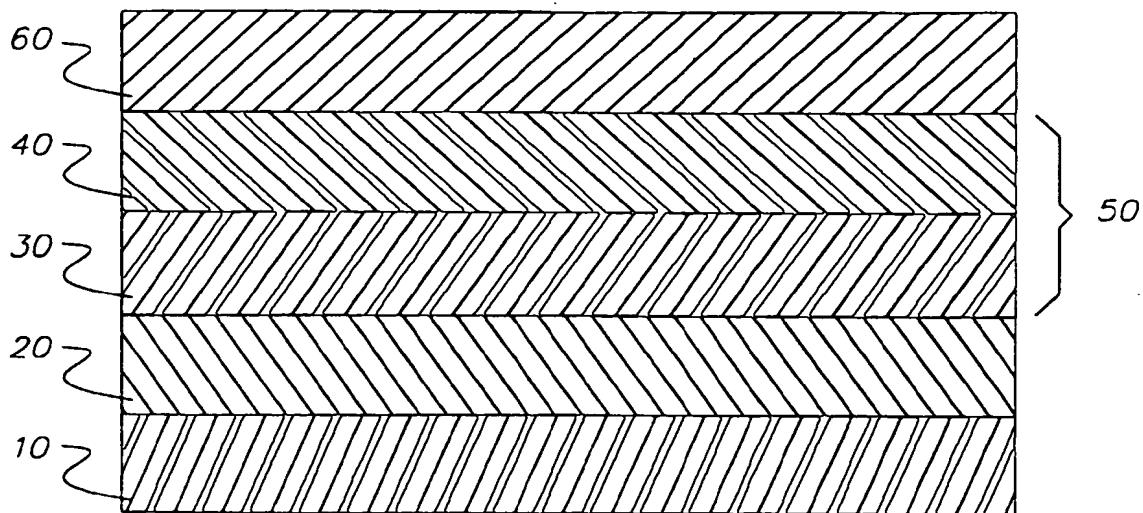


FIG. 1

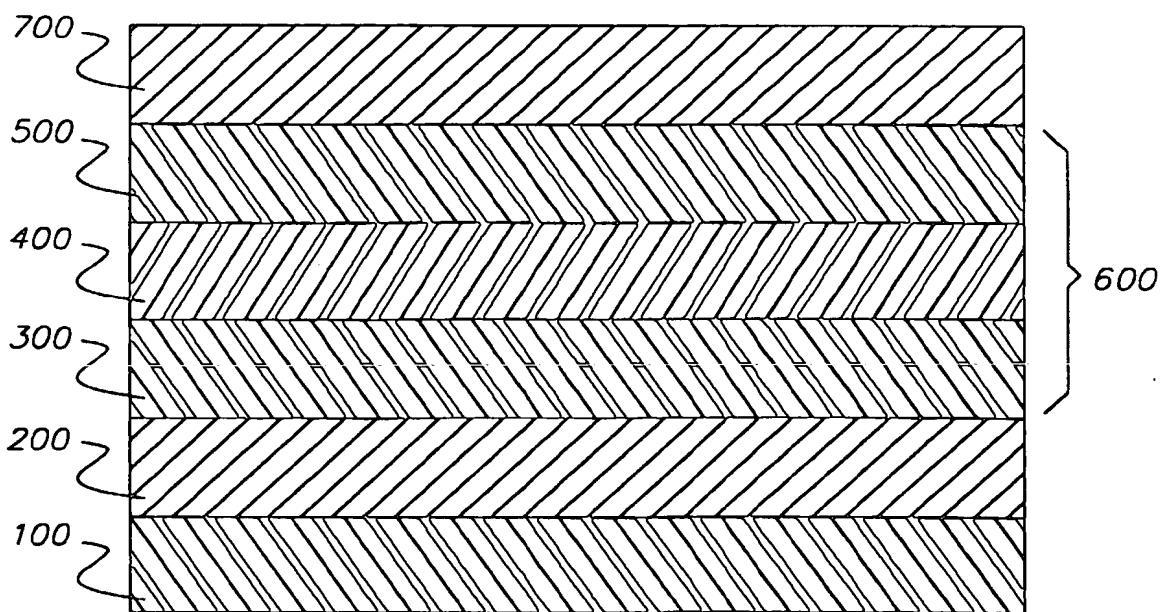


FIG. 2

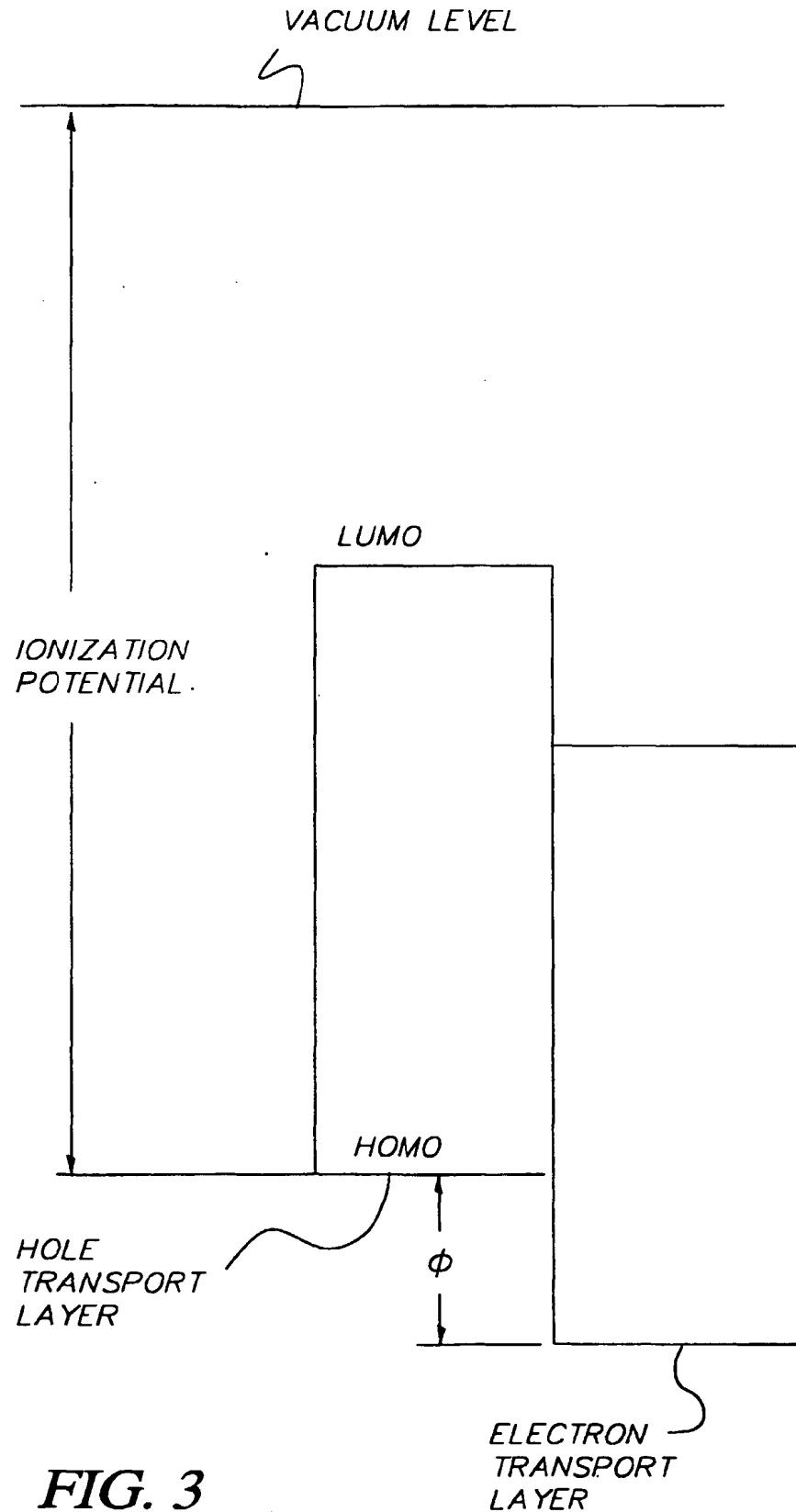
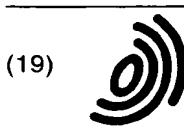


FIG. 3



(19)

Europäisches Patentamt
European Patent Office
Office européen des brevets



(11)

EP 1 009 044 A3

(12)

EUROPEAN PATENT APPLICATION

(88) Date of publication A3:
10.04.2002 Bulletin 2002/15

(51) Int Cl. 7: H01L 51/20, H05B 33/12,
H05B 33/14, H05B 33/20,
H05B 33/22, C09K 11/06,
H01L 51/30

(43) Date of publication A2:
14.06.2000 Bulletin 2000/24

(21) Application number: 99203965.1

(22) Date of filing: 25.11.1999

(84) Designated Contracting States:
AT BE CH CY DE DK ES FI FR GB GR IE IT LI LU
MC NL PT SE
Designated Extension States:
AL LT LV MK RO SI

- Tang, Ching W.
Rochester, New York 14650-2201 (US)
- Klubek, Kevin P.
Rochester, New York 14650-2201 (US)

(30) Priority: 09.12.1998 US 208172

(74) Representative:
Lewandowsky, Klaus, Dipl.-Ing. et al
Kodak Aktiengesellschaft,
Patentabteilung
70323 Stuttgart (DE)

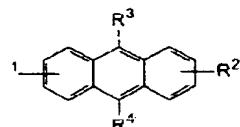
(71) Applicant: EASTMAN KODAK COMPANY
Rochester, New York 14650 (US)

(72) Inventors:

- Shi, Jianmin
Rochester, New York 14650-2201 (US)

(54) Electroluminescent device with anthracene derivatives hole transport layer

(57) An organic multilayer electroluminescent device including an anode and cathode, and including therebetween a hole transport layer; and an electron transport layer disposed in operative relationship with the hole transport layer; wherein the hole transport layer includes an organic compound having the formula:



wherein:

substituents R¹, R², R³ and R⁴ are each individually hydrogen, or alkyl of from 1 to 24 carbon atoms; aryl or substituted aryl of from 5 to 20 carbon atoms; or heteroaryl or substituted heteroaryl of from 5 to 24 carbon atoms; or fluorine, chlorine, bromine; or cyano group.

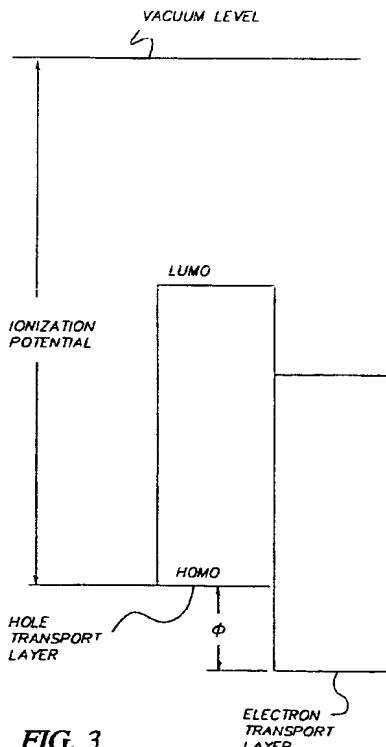


FIG. 3



European Patent
Office

EUROPEAN SEARCH REPORT

Application Number
EP 99 20 3965

DOCUMENTS CONSIDERED TO BE RELEVANT			CLASSIFICATION OF THE APPLICATION (Int.Cl.7)
Category	Citation of document with indication, where appropriate, of relevant passages	Relevant to claim	
P,X	EP 0 952 200 A (EASTMAN KODAK CO) 27 October 1999 (1999-10-27) * page 7 - page 12; claims 1-9; examples 1-6 *	1	H01L51/20 H05B33/12 H05B33/14 H05B33/20 H05B33/22 C09K11/06 H01L51/30
X	EP 0 681 019 A (TDK CORP) 8 November 1995 (1995-11-08) * page 5; claim 2 *	1	
			TECHNICAL FIELDS SEARCHED (Int.Cl.7)
			H05B C09K H01L
<p>The present search report has been drawn up for all claims</p>			
Place of search	Date of completion of the search	Examiner	
THE HAGUE	22 November 2001	Wengeler, H	
CATEGORY OF CITED DOCUMENTS			
X : particularly relevant if taken alone Y : particularly relevant if combined with another document of the same category A : technological background O : non-written disclosure P : intermediate document			
T : theory or principle underlying the invention E : earlier patent document, but published on, or after the filing date D : document cited in the application L : document cited for other reasons & : member of the same patent family, corresponding document			

**CLAIMS INCURRING FEES**

The present European patent application comprised at the time of filing more than ten claims.

- Only part of the claims have been paid within the prescribed time limit. The present European search report has been drawn up for the first ten claims and for those claims for which claims fees have been paid, namely claim(s):

1

- No claims fees have been paid within the prescribed time limit. The present European search report has been drawn up for the first ten claims.

LACK OF UNITY OF INVENTION

The Search Division considers that the present European patent application does not comply with the requirements of unity of invention and relates to several inventions or groups of inventions, namely:

see sheet B

- All further search fees have been paid within the fixed time limit. The present European search report has been drawn up for all claims.

- As all searchable claims could be searched without effort justifying an additional fee, the Search Division did not invite payment of any additional fee.

- Only part of the further search fees have been paid within the fixed time limit. The present European search report has been drawn up for those parts of the European patent application which relate to the inventions in respect of which search fees have been paid, namely claims:

- None of the further search fees have been paid within the fixed time limit. The present European search report has been drawn up for those parts of the European patent application which relate to the invention first mentioned in the claims, namely claims:

1



The Search Division considers that the present European patent application does not comply with the requirements of unity of invention and relates to several inventions or groups of inventions, namely:

1. Claim : 1

The subject-matter of claim 1 refers to a multilayer electroluminescent device comprising a hole transport layer and an electron transport layer, the hole transport layer including an anthracene based compound according to the formula of claim 1

2. Claim : 2

the hole transport layer includes an anthracene based compound according to the formula of claim 2

3. Claim : 3

the hole transport layer includes an anthracene based compound according to the formula of claim 3

4. Claim : 4

the hole transport layer includes an anthracene based compound according to the formula of claim 4

5. Claim : 5

hole transport layer including an anthracene based compound according to the formula of claim 5

6. Claim : 6

hole transport layer including an anthracene based compound according to the formula of claim 6

7. Claim : 7

hole transport layer including an anthracene based compound according to the formula of claim 7

8. Claim : 8

hole transport layer including an anthracene based compound according to the formula of claim 8

European Patent
OfficeLACK OF UNITY OF INVENTION
SHEET BApplication Number
EP 99 20 3965

The Search Division considers that the present European patent application does not comply with the requirements of unity of invention and relates to several inventions or groups of inventions, namely:

9. Claim : 9

hole transport layer including an anthracene based compound
according to the formula of claim 9

10. Claim : 10

hole transport layer including an anthracene based compound
according to the formula of claim 10

11. Claim : 11

hole transport layer including an anthracene based compound
according to the formula of claim 11

**ANNEX TO THE EUROPEAN SEARCH REPORT
ON EUROPEAN PATENT APPLICATION NO.**

EP 99 20 3965

This annex lists the patent family members relating to the patent documents cited in the above-mentioned European search report.
The members are as contained in the European Patent Office EDP file on
The European Patent Office is in no way liable for these particulars which are merely given for the purpose of information.

22-11-2001

Patent document cited in search report		Publication date		Patent family member(s)	Publication date
EP 0952200	A	27-10-1999	US	5972247 A	26-10-1999
			EP	0952200 A2	27-10-1999
			JP	11312588 A	09-11-1999
<hr/>					
EP 0681019	A	08-11-1995	DE	69511755 D1	07-10-1999
			DE	69511755 T2	13-01-2000
			EP	0681019 A2	08-11-1995
			JP	8012600 A	16-01-1996
			US	5635308 A	03-06-1997
<hr/>					

EPO FORM P0459

For more details about this annex : see Official Journal of the European Patent Office, No. 12/82

**This Page is Inserted by IFW Indexing and Scanning
Operations and is not part of the Official Record**

BEST AVAILABLE IMAGES

Defective images within this document are accurate representations of the original documents submitted by the applicant.

Defects in the images include but are not limited to the items checked:

- BLACK BORDERS**
- IMAGE CUT OFF AT TOP, BOTTOM OR SIDES**
- FADED TEXT OR DRAWING**
- BLURRED OR ILLEGIBLE TEXT OR DRAWING**
- SKEWED/SLANTED IMAGES**
- COLOR OR BLACK AND WHITE PHOTOGRAPHS**
- GRAY SCALE DOCUMENTS**
- LINES OR MARKS ON ORIGINAL DOCUMENT**
- REFERENCE(S) OR EXHIBIT(S) SUBMITTED ARE POOR QUALITY**
- OTHER:** _____

IMAGES ARE BEST AVAILABLE COPY.

As rescanning these documents will not correct the image problems checked, please do not report these problems to the IFW Image Problem Mailbox.

THIS PAGE BLANK (USPTO)